

§ 1308.47

§ 1308.47 Control of immediate precursors.

Pursuant to section 201(e) of the Act (21 U.S.C. 811(e)), the Administrator may, without regard to the findings required by subsection 201(a) or 202 (b) of the Act (21 U.S.C. 811(a) or 812(b)) and without regard to the procedures prescribed by §1308.41 or subsections 201 (a) and (b) of the Act (21 U.S.C. 811(a) and (b)), issue and publish in the FEDERAL REGISTER an order controlling an immediate precursor. The order shall designate the schedule in which the immediate precursor is to be placed, which shall be the same schedule in which the controlled substance of which it is an immediate precursor is placed or any other schedule with a higher numerical designation. An order controlling an immediate precursor shall become effective 30 days from the date of publication in the FEDERAL REGISTER, unless the Administrator finds that conditions of public health or safety necessitate an earlier effective date, in which event the Administrator shall specify in the order his findings as to such conditions.

[38 FR 8254, Mar. 30, 1973. Redesignated at 38 FR 26609, Sept. 24, 1973, and further redesignated at 62 FR 13968, Mar. 24, 1997]

§ 1308.49 Emergency scheduling.

Pursuant to 21 U.S.C. 811(h) and without regard to the requirements of 21 U.S.C. 811(b) relating to the scientific and medical evaluation of the Secretary of Health and Human Services, the Administrator may place a substance into Schedule I on a temporary basis, if he determines that such action is necessary to avoid an imminent hazard to the public safety. An order issued under this section may not be effective before the expiration of 30 days from:

(a) The date of publication by the Administrator of a notice in the FEDERAL REGISTER of his intention to issue such order and the grounds upon which such order is to be issued, and

(b) The date the Administrator has transmitted notification to the Secretary of Health and Human Services of his intention to issue such order. An order issued under this section shall be vacated upon the conclusion of a subsequent rulemaking proceeding initiated

21 CFR Ch. II (4–1–02 Edition)

under section 201(a) (21 U.S.C. 811(a)) with respect to such substance or at the end of one year from the effective date of the order scheduling the substance, except that during the pendency of proceedings under section 201(a) (21 U.S.C. 811(a)) with respect to the substance, the Administrator may extend the temporary scheduling for up to six months.

[51 FR 15318, Apr. 23, 1986. Redesignated and amended at 62 FR 13968, Mar. 24, 1997]

PART 1309—REGISTRATION OF MANUFACTURERS, DISTRIBUTORS, IMPORTERS AND EXPORTERS OF LIST I CHEMICALS

GENERAL INFORMATION

Sec.

- 1309.01 Scope of part 1309.
- 1309.02 Definitions.
- 1309.03 Information; special instructions.

FEES FOR REGISTRATION AND REREGISTRATION

- 1309.11 Fee amounts.
- 1309.12 Time and method of payment; refund.

REQUIREMENTS FOR REGISTRATION

- 1309.21 Persons required to register.
- 1309.22 Separate registration for independent activities.
- 1309.23 Separate registration for separate locations.
- 1309.24 Exemption of agents and employees.
- 1309.25 Exemption of certain controlled substance registrants.
- 1309.26 Exemption of law enforcement officials.
- 1309.27 Exemption of certain manufacturers.
- 1309.28 Exemption of distributors of regulated prescription drug products.
- 1309.29 Waiver of registration requirement for certain activities.

APPLICATION FOR REGISTRATION

- 1309.31 Time for application for registration; expiration date.
- 1309.32 Application forms; contents, signature.
- 1309.33 Filing of application; joint filings.
- 1309.34 Acceptance for filing; defective applications.
- 1309.35 Additional information.
- 1309.36 Amendments to and withdrawals of applications.

ACTION ON APPLICATIONS FOR REGISTRATION: REVOCATION OR SUSPENSION OF REGISTRATION

- 1309.41 Administrative review generally.